### **Environmental Protection Agency**

that already has a preclassifier. For example, if you use a hat-shaped preclassifier that is located immediately upstream of the probe in such a way that it forces the sample flow to change direction before entering the probe, you may not use any other preclassifier in your PM sampling system.

(2) Other components. You may request to use other PM conditioning components upstream of a PM preclassifier, such as components that condition humidity or remove gaseousphase hydrocarbons from the diluted exhaust stream. You may use such components only if we approve them under §1065.10.

#### § 1065.150 Continuous sampling.

You may use continuous sampling techniques for measurements that involve raw or dilute sampling. Make sure continuous sampling systems meet the specifications in §1065.145. Make sure continuous analyzers meet the specifications in subparts C and D of this part.

## § 1065.170 Batch sampling for gaseous and PM constituents.

Batch sampling involves collecting and storing emissions for later analysis. Examples of batch sampling include collecting and storing gaseous emissions in a bag and collecting and storing PM on a filter. You may use batch sampling to store emissions that have been diluted at least once in some way, such as with CVS, PFD, or BMD. You may use batch-sampling to store undiluted emissions only if we approve

it as an alternate procedure under §1065.10.

(a) Sampling methods. For batch sampling, extract the sample at a rate proportional to the exhaust flow. If you extract from a constant-volume flow rate, sample at a constant-volume flow rate. If you extract from a varying flow rate, vary the sample rate in proportion to the varying flow rate. Validate proportional sampling after an emission test as described in §1065.545. Use storage media that do not change measured emission levels (either up or down). For example, do not use sample bags for storing emissions if the bags are permeable with respect to emissions or if they off-gas emissions. As another example, do not use PM filters that irreversibly absorb or adsorb gases.

(b) Gaseous sample storage media. Store gas volumes in sufficiently clean containers that minimally off-gas or allow permeation of gases. Use good engineering judgment to determine acceptable thresholds of storage media cleanliness and permeation. To clean a container, you may repeatedly purge and evacuate a container and you may heat it. Use a flexible container (such as a bag) within a temperature-controlled environment, or use a temperature controlled rigid container that is initially evacuated or has a volume that can be displaced, such as a piston and cylinder arrangement. Use containers meeting the specifications in the following table, noting that you may request to use other container materials under §1065.10:

TABLE 1 OF § 1065.170—GASEOUS BATCH SAMPLING CONTAINER MATERIALS

Emissions	Engines	
	Compression-ignition, two-stroke spark ignition, 4-stroke spark-ignition <19 kW	All other engines
CO, CO <sub>2</sub> , O <sub>2</sub> , CH <sub>4</sub> , C <sub>2</sub> H <sub>6</sub> , C <sub>3</sub> H <sub>8</sub> , NO, NO <sub>2</sub> <sup>1</sup> .	Tedlar <sup>TM</sup> , <sup>2</sup> Kynar <sup>TM</sup> , <sup>2</sup> Teflon <sup>TM</sup> , <sup>3</sup> or 300 series stainless steel <sup>3</sup> .	Tedlar <sup>TM</sup> , <sup>2</sup> Kynar <sup>TM</sup> , <sup>2</sup> Teflon <sup>TM</sup> , <sup>3</sup> or 300 series stainless steel <sup>3</sup>
THC, NMHC	Teflon™4 or 300 series stainless steel <sup>4</sup>	Tedlar <sup>™</sup> , <sup>2</sup> Kynar <sup>™</sup> , <sup>2</sup> Tef- lon <sup>™</sup> , <sup>3</sup> or 300 series stain- less steel <sup>3</sup>

<sup>&</sup>lt;sup>1</sup> As long as you prevent aqueous condensation in storage container.

<sup>&</sup>lt;sup>2</sup>Up to 40 °C.

<sup>&</sup>lt;sup>3</sup>Up to 202 °C.

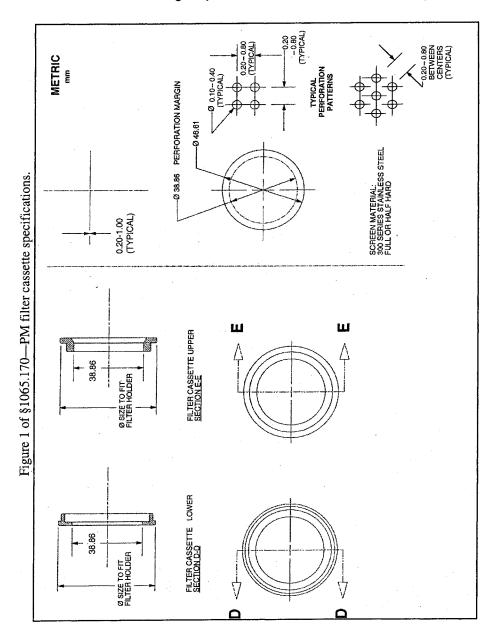
<sup>&</sup>lt;sup>4</sup>At (191 ±11) °C

### § 1065.170

- (c) *PM sample media*. Apply the following methods for sampling particulate emissions:
- (1) If you use filter-based sampling media to extract and store PM for measurement, your procedure must meet the following specifications:
- (i) If you expect that a filter's total surface concentration of PM will exceed 0.473 mm/mm² for a given test interval, you may use filter media with a minimum initial collection efficiency of 98%; otherwise you must use a filter media with a minimum initial collection efficiency of 99.7%. Collection efficiency must be measured as described in ASTM D 2986-95a (incorporated by reference in §1065.1010), though you may rely on the sample-media manufacturer's measurements reflected in their product ratings to show that you meet applicable requirements.
- (ii) The filter must be circular, with an overall diameter of  $46.50 \pm 0.6$  mm and an exposed diameter of at least 38 mm. See the cassette specifications in paragraph (c)(1)(vi) of this section.
- (iii) We highly recommend that you use a pure PTFE filter material that does not have any flow-through support bonded to the back and has an overall thickness of 40 ±20 μm. An inert polymer ring may be bonded to the periphery of the filter material for support and for sealing between the filter cassette parts. We consider Polymethylpentene (PMP) and PTFE inert materials for a support ring, but other inert materials may be used. See the cassette specifications in paragraph (c)(1)(v) of this section. We allow the use of PTFE-coated glass fiber filter material, as long as this filter media selection does not affect your ability to demonstrate compliance with the applicable standards, which we base on a pure PTFE filter material. Note that we will use pure PTFE filter material for compliance testing, and we may require you to use pure

PTFE filter material for any compliance testing we require, such as for selective enforcement audits.

- (iv) You may request to use other filter materials or sizes under the provisions of § 1065.10.
- (v) To minimize turbulent deposition and to deposit PM evenly on a filter, use a  $12.5^{\circ}$  (from center) divergent cone angle to transition from the transferline inside diameter to the exposed diameter of the filter face. Use 300 series stainless steel for this transition.
- (vi) Maintain sample velocity at the filter face at or below 100 cm/s, where filter face velocity is the measured volumetric flow rate of the sample at the pressure and temperature upstream of the filter face, divided by the filter's exposed area.
- (vii) Use a clean cassette designed to the specifications of Figure 1 of §1065.170 and made of any of the following materials: Delrin<sup>TM</sup>, 300 series stainless steel, polycarbonate, acrylonitrile-butadiene-styrene (ABS) resin, or conductive polypropylene. We recommend that you keep filter cassettes clean by periodically washing or wiping them with a compatible solvent applied using a lint-free cloth. Depending upon cassette material. ethanol vour (C<sub>2</sub>H<sub>5</sub>OH) might be an acceptable solvent. Your cleaning frequency will depend on your engine's PM and HC emissions.
- (viii) If you store filters in cassettes in an automatic PM sampler, cover or seal individual filter cassettes after sampling to prevent communication of semi-volatile matter from one filter to another.
- (2) You may use other PM sample media that we approve under §1065.10, including non-filtering techniques. For example, you might deposit PM on an inert substrate that collects PM using electrostatic, thermophoresis, inertia, diffusion, or some other deposition mechanism, as approved.



# § 1065.190 PM-stabilization and weighing environments for gravimetric analysis.

(a) This section describes the two environments required to stabilize and

weigh PM for gravimetric analysis: the PM stabilization environment, where filters are stored before weighing; and the weighing environment, where the balance is located. The two environments may share a common space.